

GenCore version 5.1.4_p5_4578
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OM protein - protein search, using sw model

Run on: March 14, 2003, 17:35:22 ; Search time 43 seconds
(without alignments)
3210.410 Million cell updates/sec

Title: US-09-887-527A-60

Perfect score: 5952

Sequence: 1 MYLVAGDRGLAGCGHLLVSL.....GFYSMQKNHLQADNFYQTV 1036

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 908470 seqs, 133250620 residues

Total number of hits satisfying chosen parameters: 908470

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : A_Geneseq_101002.*

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- 2: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA1981.DAT.*
- 3: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA1982.DAT.*
- 4: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA1983.DAT.*
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- 18: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA1997.DAT.*
- 19: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA1998.DAT.*
- 20: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA1999.DAT.*
- 21: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2000.DAT.*
- 22: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2001.DAT.*
- 23: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2002.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	5952	100.0	1036	AAE18852	Human pharmaceutical
2	5906	99.2	1036	AAI82776	Human chordin rela
3	5906	99.2	1036	AAI53034	Human secreted pro
4	5906	99.2	1036	AAU12242	Human PRO4330 poly
5	5901	99.1	1036	AAU07141	Human CRIM1 protei
6	5402.5	90.8	1037	AAU07142	Mouse CRIM1 protei
7	4969	83.5	1048	AAU07143	Chicken CRIM1 prot
8	4167	70.0	732	AAI61140	Human NOV10 protei
9	2828.5	47.5	503	ABG66681	Human novel polype
10	2254	37.9	400	AAI82775	Human chordin rela

11	2254	37.9	400	21	AAI53033	Human secreted pro
12	1575	26.5	322	21	AAI40954	Human ORFX ORF18
13	1307	22.0	872	22	AAU07149	C. elegans CRIM1 p
14	1042	17.5	193	22	AAI25238	Human protein sequ
15	582	9.8	225	23	AAU83112	Novel secreted pro
16	506	8.5	810	18	AAI37500	Human nel-related
17	429.5	7.2	816	18	AAI37501	Human nel-related
18	419.5	7.0	361	22	AAI99918	Human polypeptide
19	419	7.0	72	22	ABR28970	Peptide #1621 enco
20	419	7.0	72	22	ABR34137	Peptide #1643 enco
21	419	7.0	72	22	ABR19578	Protein #1577 enco
22	419	7.0	72	22	AAI54928	Human brain expres
23	419	7.0	72	22	AAI67308	Human bone marrow
24	419	7.0	72	22	AAI15145	Peptide #1579 enco
25	419	7.0	72	22	AAI27600	Peptide #1637 enco
26	419	7.0	72	22	AAI02886	Peptide #1568 enco
27	419	7.0	72	23	ABG36955	Human peptide enco
28	416	7.0	73	22	ABR34229	Peptide #1735 enco
29	416	7.0	73	22	ABR19665	Protein #1664 enco
30	416	7.0	73	22	AAI55023	Human brain expres
31	416	7.0	73	22	AAI67408	Human bone marrow
32	416	7.0	73	22	AAI15239	Peptide #1673 enco
33	416	7.0	73	22	AAI27700	Peptide #1737 enco
34	416	7.0	73	22	AAI02982	Peptide #1664 enco
35	416	7.0	73	23	ABG37035	Human peptide enco
36	413.5	6.9	2444	23	ABR07821	Constitutively act
37	413.5	6.9	3680	22	ABR70878	Drosophila melanog
38	404	6.8	445	22	AAE07062	Human gene 12 enco
39	404	6.8	445	23	ABG65086	Human albumin fus1
40	404	6.8	464	22	AAI07119	Human gene 12 enco
41	402	6.8	685	23	AAI99292	Human chordin-like
42	394	6.6	2809	23	AAI66169	Human fibrillin 3
43	378.5	6.4	2912	22	ABG06402	Novel human diagno
44	378	6.4	627	23	AAI99293	Human chordin-like
45	374	6.3	63	22	ABR29742	Peptide #2393 enco

ALIGNMENTS

RESULT 1
AAE18852
ID AAE18852 standard; Protein; 1036 AA.
XX AAE18852;
AC AAE18852;
DT 17-MAY-2002 (first entry)
XX Human pharmaceutical compound protein for cancer treatment.
XX Human; pharmaceutical composition; compound I; tumour; psoriasis; cancer;
XX rheumatoid arthritis; vascular endothelial growth factor; VEGF; therapy;
XX neovascular glaucoma; compound II; angiotensin/Tie receptor system;
XX retinopathy; glomerulonephritis; diabetic nephropathy; nephrosclerosis;
XX thrombotic microangiopathic syndrome; transplantation; glomerulopathy;
XX fibrotic disease; cirrhotic liver; proliferative disease; nephropathy;
XX ophthalmological; arteriosclerosis; cytotatic; hepatotropic; oedema.
XX Homo sapiens.
XX WO200197850-A2.
XX 27-DEC-2001.
XX 20-JUN-2001; 2001WO-EP06976.
XX 23-JUN-2000; 2000EP-0250194.
XX 28-JUN-2000; 2000EP-0250214.
XX (SCHD) SCHERING AG.
XX (SIEM/) SIEMEISTER G.
XX (HABE/) HABEREY M.
XX (THIE/) THIERAUCH K.

XX Siemeister G, Haberey M, Thierauch K;
XX WPI; 2002-179543/23.
DR N-PSDB; AAD29965.
XX Novel composition useful for treating cancer, comprises agents
PT interfering with vascular endothelial growth factor/VEGF receptor
PT system activity and agents interfering with Angiopoietin/Tie receptor
PT system function
XX Claim 13; Page 69-70; 79pp; English.
XX The present invention relates to a pharmaceutical composition comprising
CC a combination of substances (compound I) interfering with the biological
CC activity of vascular endothelial growth factor (VEGF)/VEGF receptor
CC systems and substances (compound II) interfering with the biological
CC function of angiopoietin/Tie receptor systems. The pharmaceutical
CC composition is useful for the production of a medicament for the
CC treatment of tumours, cancers, psoriasis, arthritis, such as rheumatoid
CC arthritis, haemangioma, angiofibroma, eye disease such as diabetic
CC retinopathy, neovascular glaucoma, kidney disease such as
CC glomerulonephritis, diabetic nephropathy, malignant nephrosclerosis,
CC thrombotic microangiopathic syndrome, transplantation rejections and
CC glomerulopathy, fibrotic disease such as cirrhotic liver, mesangial cell
CC proliferative diseases, arteriosclerosis, damage of nerve tissues,
CC suppression of the ascites formation in patients and suppression of VEGF
CC oedemas. The present sequence is human pharmaceutical compound protein
CC used in the invention.
XX Sequence 1036 AA;
SQ
Query Match 100.0%; Score 5952; DB 23; Length 1036;
Best Local Similarity 100.0%; Pred. No. 0;
Matches 1036; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 MYLVAGDRGLAGCHLLVSLGLLLPARSGTRALVCLPCDESKCEPRNRPGSVQVGC 60
Db 1 MYLVAGDRGLAGCHLLVSLGLLLPARSGTRALVCLPCDESKCEPRNRPGSVQVGC 60
QY 61 GCCTCASQGNESCGGTGFIYGTCDRLGRCVIRPPLNGDSLTYEAGVCEDENTDQLL 120
Db 61 GCCTCASQGNESCGGTGFIYGTCDRLGRCVIRPPLNGDSLTYEAGVCEDENTDQLL 120
QY 121 GFKPCNENLIAGNIINGKCECNIITRTCSNPFEPFQDMCLSAKRIEIEKPDCKSARCE 180
Db 121 GFKPCNENLIAGNIINGKCECNIITRTCSNPFEPFQDMCLSAKRIEIEKPDCKSARCE 180
QY 181 VQSPRCPEDSVLEIYAPGECCLPLSRVCVNPAGCLRKVCQPGNLIIVSKASKGPE 240
Db 181 VQSPRCPEDSVLEIYAPGECCLPLSRVCVNPAGCLRKVCQPGNLIIVSKASKGPE 240
QY 241 CCCLYECKPVGVDCRTVECPVQQTACPPDSYETOVRLTADGCCPLTRCCLSLGCGF 300
Db 241 CCCLYECKPVGVDCRTVECPVQQTACPPDSYETOVRLTADGCCPLTRCCLSLGCGF 300
QY 301 PVCEVSGTPRIVSRGDTGPKCCDVFECVNDTKPACVFNNVEYDGMDFMDCNRCFCRCQ 360
Db 301 PVCEVSGTPRIVSRGDTGPKCCDVFECVNDTKPACVFNNVEYDGMDFMDCNRCFCRCQ 360
QY 361 GGVAICFTACQGEINERYVYVPEGCCPCVEDPVYFPNPNAGCYANGLILAHGDRWRDD 420
Db 361 GGVAICFTACQGEINERYVYVPEGCCPCVEDPVYFPNPNAGCYANGLILAHGDRWRDD 420
QY 421 CTCQCQVNGRHCVAATVCGTCTNPKVPGECPCVCEEPTIITVDPACGELSNCNLTTRK 480
Db 421 CTCQCQVNGRHCVAATVCGTCTNPKVPGECPCVCEEPTIITVDPACGELSNCNLTTRK 480
QY 481 DCINGFKRDHNGCRTCCINTQELCSERKQGGTLCNPGFGLTDAQNCICECPRPKRCR 540
Db 481 DCINGFKRDHNGCRTCCINTQELCSERKQGGTLCNPGFGLTDAQNCICECPRPKRCR 540
QY 541 PIICDKYCPILGLLNKNGCDICRCKPELSCSKICPLGFOQDSHGCLICKRASASAG 600

Db 541 PIICDKYCPILGLLNKNGCDICRCKPELSCSKICPLGFOQDSHGCLICKRASASAG 600
QY 601 PPIISGTCCLTVDGHKNEESWHDGCRCYCLNGREMCALITCPVPACGNPTIHPGQCCP 660
Db 601 PPIISGTCCLTVDGHKNEESWHDGCRCYCLNGREMCALITCPVPACGNPTIHPGQCCP 660
QY 661 SCADDFVQVKPELSTPSCIPAGGGEYFVEGETWNIDSTQCTCHSGRVLCEVCPPLLC 720
Db 661 SCADDFVQVKPELSTPSCIPAGGGEYFVEGETWNIDSTQCTCHSGRVLCEVCPPLLC 720
QY 721 QNPSTQSCCPQCTDQPFPSLRNNSVPNYCKNDEGDIPLAAESWKPVDVCTSCICIDS 780
Db 721 QNPSTQSCCPQCTDQPFPSLRNNSVPNYCKNDEGDIPLAAESWKPVDVCTSCICIDS 780
QY 781 VTSCSESCPSVSCERPVLRKGQCCPYCIKDTIPKVVCHFSGRAYADEERWDLDSCTHC 840
Db 781 VTSCSESCPSVSCERPVLRKGQCCPYCIKDTIPKVVCHFSGRAYADEERWDLDSCTHC 840
QY 841 YCLOQTLCSTVSCPPPCVPEPINVEGSCCPMCPMYPPEPTNPIEKTNRHGEVDLEVP 900
Db 841 YCLOQTLCSTVSCPPPCVPEPINVEGSCCPMCPMYPPEPTNPIEKTNRHGEVDLEVP 900
QY 901 LWPTSENDIVLPRDMGHQLQVDRNRLHPSESSLDSTASVVVPIIICLSIIIAFLFI 960
Db 901 LWPTSENDIVLPRDMGHQLQVDRNRLHPSESSLDSTASVVVPIIICLSIIIAFLFI 960
QY 961 NOKKQWIPLLCWYRTPTKPSLNNQLVSDCKKGTTRVQVDSQRMRLIAEPDARFSGFYS 1020
Db 961 NOKKQWIPLLCWYRTPTKPSLNNQLVSDCKKGTTRVQVDSQRMRLIAEPDARFSGFYS 1020
QY 1021 MOKNHLQADNFYQTV 1036
Db 1021 MOKNHLQADNFYQTV 1036
RESULT 2
AAY82776
ID AAY82776 standard; Protein; 1036 AA.
XX AC AAY82776;
XX DT 19-JUN-2000 (first entry)
XX Human chordin related protein (Clone dj191_19).
KW Chordin related protein; cartilage; bone; connective tissue;
KW periodontal disease; osteoporosis; burn; incision; ulcer; neuron;
KW astrocyte; glial cell; transplantation; nerve; epidermis; muscle;
KW liver; brain; lung; cardiac; pancreas; kidney; growth;
KW differentiation; TGF-Beta; angiogenesis; chemotaxis;
KW chemoattraction; collagen synthesis; fibrosis; cell adhesion;
KW cell migration; fertility; reproduction; haematopoiesis;
KW erythroid cell; tumour; dietary supplement; growth medium.
OS Homo sapiens.
XX WO200009551-A1.
XX 24-FEB-2000.
XX 10-AUG-1999; 99WO-US18117.
XX 10-AUG-1998; 98US-0095880.
XX 06-MAY-1999; 99US-0306111.
XX (GEMY) GENETICS INST INC.
XX Jacobs K, McCoy JM, Lavallie ER, Collins-racie LA, Merberg D;
PI Treacy M, Diblasio-smith E, Widom A;
XX WPI; 2000-205978/18.
DR N-PSDB; AA293172.

XX New polynucleotides encoding secreted human proteins, useful for
PT treating e.g. broken bones, craniofacial defects, periodontal disease,
PT osteoporosis, burns, incisions or ulcers
XX
PS Claim 21; Page 94-98; 105pp; English.
XX
CC The human chordin related protein and polynucleotides encoding them
CC are predicted to have biological activities which would make them
CC suitable for treating, preventing or ameliorating medical conditions
CC which involve defects in cartilage, bone or connective tissue
CC formation and damage to cartilage, bone or connective tissue, e.g.
CC broken bones, congenital, trauma-induced, or
CC oncologic-resection-induced craniofacial defects, periodontal
CC disease, defects in the periodontal ligament or attachment apparatus,
CC damage to the periodontal ligament or attachment apparatus,
CC osteoporosis, burns, incisions or ulcers. The proteins may also
CC affect neuronal, astrocytic, and glial cell survival and therefore be
CC useful in transplantation and treatment of conditions exhibiting a
CC decrease in neuronal survival and repair. The proteins may also be
CC useful for the treatment of conditions related to other types of
CC tissue, such as nerve, epidermis, muscle, and other organs such as
CC liver, brain, lung, cardiac, pancreas, and kidney tissue. The
CC proteins may further be useful for the treatment of relatively
CC undifferentiated cell populations, such as embryonic cells, or stem
CC cells, to enhance growth and/or differentiation of the cells.
CC The proteins may also have other useful properties characteristic of
CC the TGF-beta superfamily of proteins. Such properties include
CC angiogenic, chemotactic, and/or chemoattractant properties, and
CC effects on cells including induction or inhibition of collagen
CC synthesis, fibrosis, differentiation responses, cell proliferative
CC responses, and responses involving cell adhesion, migration, and
CC extracellular matrices. These properties make the proteins potential
CC agents for wound healing, reduction of fibrosis, and reduction of
CC scar tissue formation. Chordin-related proteins may also be useful
CC for advancement of the onset of fertility in sexually immature
CC mammals, so as to increase the lifetime reproductive performance of
CC domestic animals such as cows, sheep and pigs. Chordin-related
CC proteins may also be useful in modulating hematopoiesis by inducing
CC the differentiation of erythroid cells, for suppressing the
CC development of gonadal tumors, or for augmenting the activity of
CC BMPs. The proteins may also have value as a dietary supplement, or
CC as a component of cell culture media.
XX
SQ Sequence 1036 AA;

Query Match 99.2%; Score 5906; DB 21; Length 1036;
Best Local Similarity 99.3%; Pred. No. 0;
Matches 1029; Conservative 2; Mismatches 5; Indels 0; Gaps 0;

QY 1 MYLVAGDRGLAGCGHLLVSLGLLLPARSGTRALVCLPCDESKCEPRNRPGSIVQVC 60
DB 1 MYLVAGDRGLAGCGHLLVSLGLLLPARSGTRALVCLPCDESKCEPRNRPGSIVQVC 60
QY 61 GCCTCASQNBSCGGTGIYGTCDRLGRCVIRPPLNGDSLTYEAGVCEDENTDQLL 120
DB 61 GCCTCASQNBSCGGTGIYGTCDRLGRCVIRPPLNGDSLTYEAGVCEDENTDQLL 120
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DB 121 GFKPCNEMLIAGCNLIINGKCECNRITCSNPFEPFSDQMCLSALKRTEERKPDCKARCE 180
QY 181 VQFSRCPEDSVLIEGYAPPECCPLPSRCVNCNAGCLRKVCQPCNLIIVSKASGRPGE 240
DB 181 VQFSRCPEDSVLIEGYAPPECCPLPSRCVNCNAGCLRKVCQPCNLIIVSKASGRPGE 240
QY 241 CCCLYECKPVGVDCTVECPVQQTACPPDPDSYETQVRLTADGGCTLPTRCECLSLGCGF 300
DB 241 CCCLYECKPVGVDCTVECPVQQTACPPDPDSYETQVRLTADGGCTLPTRCECLSLGCGF 300
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DB 301 PVCEVGSTPRIVSRGDTGPGCCDFECVNDTKPACVFNVEYDGMFRMDNCRFCRCQ 360

QY 361 GGVAICTAOCGEINCERYVYPGECCPVCEDPVYPFNNPAGCYANGLILAHGDRWREDD 420
DB 361 GGVAICTAOCGEINCERYVYPGECCPVCEDPVYPFNNPAGCYANGLILAHGDRWREDD 420
QY 421 CTFCQCVNGERHCVATVCGQTCCTNPVKVPGCECPVCEPTIITVDPAGGELSNCITLTK 480
DB 421 CTFCQCVNGERHCVATVCGQTCCTNPVKVPGCECPVCEPTIITVDPAGGELSNCITLTK 480
QY 481 DCINGFKRDHNGRTQCIINTQELCSERKOGCTLNCFFGLTDAQNCETCECPRPKCR 540
DB 481 DCINGFKRDHNGRTQCIINTQELCSERKOGCTLNCFFGLTDAQNCETCECPRPKCR 540
QY 541 PIICDKYCPGLGLKNKHGCDICRCKPELSCSKICPLGFQODSHGCLICKCREASASAG 600
DB 541 PIICDKYCPGLGLKNKHGCDICRCKPELSCSKICPLGFQODSHGCLICKCREASASAG 600
QY 601 PPIILSGTCLTVDGHGHHKNEESHDGRCYCLNGREWCALITCPVPACGNPTIHPGCCP 660
DB 601 PPIILSGTCLTVDGHGHHKNEESHDGRCYCLNGREWCALITCPVPACGNPTIHPGCCP 660
QY 661 SCADDFVQKPELSTPSICHAPGGYFVEGETWNIDSTQCTCHSGRVLCTEVCPPLLC 720
DB 661 SCADDFVQKPELSTPSICHAPGGYFVEGETWNIDSTQCTCHSGRVLCTEVCPPLLC 720
QY 721 QNPSRTQDSCCPOCTDQPPRPSLSRNNSVNYCKNDEGDIPLAESWKPDVCTSCICIDS 780
DB 721 QNPSRTQDSCCPOCTDQPPRPSLSRNNSVNYCKNDEGDIPLAESWKPDVCTSCICIDS 780
QY 781 VISCFSESCPSVSCERPVLRKQCCPCYIKDTIPKVVCHFSKAYADEERWDLDSCTHC 840
DB 781 VISCFSESCPSVSCERPVLRKQCCPCYIKDTIPKVVCHFSKAYADEERWDLDSCTHC 840
QY 841 YCLOGQTLCTSVCCPLPCVEPINVSGCCPMCPMVVPEPTNIPIEKNHREGVLEVP 900
DB 841 YCLOGQTLCTSVCCPLPCVEPINVSGCCPMCPMVVPEPTNIPIEKNHREGVLEVP 900
QY 901 LWPTPSNDIVHLPRDMGHQVQDYRDNRLHPSDSSLSIASVVPFIIICLSIIIAFLFI 960
DB 901 LWPTPSNDIVHLPRDMGHQVQDYRDNRLHPSDSSLSIASVVPFIIICLSIIIAFLFI 960
QY 961 NQKQWIPLLCWYRTPTKPSLNNQLVSDCKKGTQVQVDSQRMRLRIAPDARFSGFYS 1020
DB 961 NQKQWIPLLCWYRTPTKPSLNNQLVSDCKKGTQVQVDSQRMRLRIAPDARFSGFYS 1020
QY 1021 MQKQNHLOADNFQTV 1036
DB 1021 MQKQNHLOADNFQTV 1036

RESULT 3
AAV53034
ID AAV53034 standard; Protein; 1036 AA.
XX
AC AAV53034;
XX
DT 29-FEB-2000 (first entry)
XX
DE Human secreted protein clone dj167_19 protein sequence SEQ ID NO:74.
XX
KW Human; secreted protein; nutritional; cytokine; cell proliferation;
KW differentiation; immune stimulating; vaccine; suppression;
KW haematopoiesis regulation; tissue growth; activin; inhibin;
KW chemotactic; chemokinetic; haemostatic; thrombolytic; receptor;
KW ligand; anti-inflammatory; cadherin; tumour invasion suppressor;
KW tumour inhibition; gene therapy.
XX
OS Homo sapiens.
XX
PN WO957132-A1.
XX
PD 11-NOV-1999.
XX

PF	07-MAY-1999;	99WO-US09970.	
XX			
PR	07-MAY-1998;	98US-0084564.	
PR	02-JUN-1998;	98US-0087645.	
PR	22-JUL-1998;	98US-0093712.	
PR	31-JUL-1998;	98US-0094935.	
PR	10-AUG-1998;	98US-0095880.	
PR	11-AUG-1998;	98US-0096068.	
PR	06-MAY-1999;	99US-0096068.	
XX			
PA	(GEM) GENETICS INST INC.		
XX			
PI	Jacobs K, McCoy JM, Lavallie ER, Collins-Racie LA, Evans C;		
PI	Merberg D, Treacy M, Agostino MJ, Steininger RJ, Bowman MR;		
PI	DiBlasio-Smith E, Widom A;		
XX			
DR	WPI: 2000-052937/04.		
DR	N-PSDB; AA233352.		
XX			
PT	New polynucleotides encoding secreted human proteins, derived from		
PT	adult placenta, adult retina, fetal brain, fetal -		
XX			
PS	Claim 83; Page 426-429; 492pp; English.		
XX			
CC	The present invention describes new human secreted proteins which were		
CC	isolated from adult placenta, adult retina, foetal brain, foetal kidney,		
CC	adult blood, adult brain, adult thyroid, adult bladder, adult neural		
CC	tissue, adult testes, and adult lymph node cDNA libraries. The human		
CC	secreted proteins, and the polynucleotides encoding them, are predicted		
CC	to have biological activities which would make them suitable for		
CC	treating, preventing or ameliorating medical conditions in humans and		
CC	animals. Suggested activities include nutritional activity, cytokine		
CC	and cell proliferation/differentiation activity, immune stimulating		
CC	(e.g. as vaccines) or suppressing activity, haematopoiesis regulating		
CC	activity, tissue growth activity, activin/inhibin activity,		
CC	chemotactic/chemokinetic activity, haemostatic and thrombolytic		
CC	activity, receptor/ligand activity, anti-inflammatory activity,		
CC	cacharin/tumour invasion suppressor activity, and tumour inhibition		
CC	activity. The polynucleotides are also stated to be useful for gene		
CC	therapy. AA233316 to AA233373 encode human secreted proteins, and		
CC	AA252998 to AA253060 represent human secreted proteins, given in the		
CC	present invention.		
XX			
SQ	Sequence 1036 AA;		
	Query Match 99.2%; Score 5906; DB 21; Length 1036;		
	Best Local Similarity 99.3%; Pred. No. 0;		
	Matches 1029; Conservative 2; Mismatches 5; Indels 0; Gaps 0;		
QY	1 MYLVAGDRGLAGGCHLLVSLGLLLPARSGTRALVCLPCDESKCEEPNRCPSIVQGYC 60		
DB	1 MYLVAGDRGLAGGCHLLVSLGLLLPARSGTRALVCLPCDESKCEEPNRCPSIVQGYC 60		
QY	61 GCCTCASOGNESCGETFGTGCORGLRCVIRPPLNGDSLTFEYAGVCEDENWTDOLL 120		
DB	61 GCCTCASOGNESCGETFGTGCORGLRCVIRPPLNGDSLTFEYAGVCEDENWTDOLL 120		
QY	121 GFKPCNENIAGNIINGKCECNIITRTCSNPFEFFSQDMCLSAKRIEKEKPDCKSKARCE 180		
DB	121 GFKPCNENIAGNIINGKCECNIITRTCSNPFEFFSQDMCLSAKRIEKEKPDCKSKARCE 180		
QY	181 VQSPRCPEDSVLIEGYPGECCLPSRCVGNPAGCLRKVCQPNLTLVSKASKGKPE 240		
DB	181 VQSPRCPEDSVLIEGYPGECCLPSRCVGNPAGCLRKVCQPNLTLVSKASKGKPE 240		
QY	241 CCLYLECKPVFVGVDCTVPCPTVQQTACPPDSYETQVRLTADGCCTLPTRCECLSLCGF 300		
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QY	301 PVCVGSTPRIVSRGDTGTPGKCCDFECVNDTKPACVFNNVEYDGMFRMDNCRFCRCQ 360		
DB	301 PVCVGSTPRIVSRGDTGTPGKCCDFECVNDTKPACVFNNVEYDGMFRMDNCRFCRCQ 360		

Qy	361	GGVAICFTTAQCGEINCERYVYPGECCPVCEDPVYPNNPAGCYANGLIIAHDWRREDD	420
Db	361	GGVAICFTTAQCGEINCERYVYPGECCPVCEDPVYPNNPAGCYANGLIIAHDWRREDD	420
Qy	421	CTFCQCVNGERHCVATVCGQTNPVAVPGECPCVCEPTIITVDPPACGELSNCITLTKR	480
Db	421	CTFCQCVNGERHCVATVCGQTNPVAVPGECPCVCEPTIITVDPPACGELSNCITLTKR	480
Qy	481	DCINGFKRDHNGCRCTQCINTQELCSERKOGCTLNCRCFGLTDAQNCCEICECPRPKCR	540
Db	481	DCINGFKRDHNGCRCTQCINTQELCSERKOGCTLNCRCFGLTDAQNCCEICECPRPKCR	540
Qy	541	PIICDKYCPGLLKNKHGCDICRCKKPELSCSKICPLGFQDDSHGCLICKREASASAG	600
Db	541	PIICDKYCPGLLKNKHGCDICRCKKPELSCSKICPLGFQDDSHGCLICKREASASAG	600
Qy	601	PPILSGTCLTVDGHGHHNEESWHDGCRECYCLNGREMCALITCPVACGNPTIHPGQCCP	660
Db	601	PPILSGTCLTVDGHGHHNEESWHDGCRECYCLNGREMCALITCPVACGNPTIHPGQCCP	660
Qy	661	SCADDFVVKPELSTPSICHAPGGYFVEGETWIDISCTQCTCHSGRVLCETEVCPPLLC	720
Db	661	SCADDFVVKPELSTPSICHAPGGYFVEGETWIDISCTQCTCHSGRVLCETEVCPPLLC	720
Qy	721	QNPRTQDSCCPQCTDQFPFRSLRNNSVNYCKNDEGDIPLAAESWKPDVCTSCICIDS	780
Db	721	QNPRTQDSCCPQCTDQFPFRSLRNNSVNYCKNDEGDIPLAAESWKPDVCTSCICIDS	780
Qy	781	VISCFSESCPSVSCERPVLRKGCCPYCIKDTIPKVVCHFSCKAYADEERWDLDSCTHC	840
Db	781	VISCFSESCPSVSCERPVLRKGCCPYCIKDTIPKVVCHFSCKAYADEERWDLDSCTHC	840
Qy	841	YCLOGQTLCTSVSCPPPLPCVPEPINVEGSCCPMCEMYPVEPTNPIEKTNRHGEVDLEVP	900
Db	841	YCLOGQTLCTSVSCPPPLPCVPEPINVEGSCCPMCEMYPVEPTNPIEKTNRHGEVDLEVP	900
Qy	901	LWPTPSENDIVHLPRDMGHLQVDYRDLNRLHPSDESSLDSTASVVVPIIICLSIIIAFLFI	960
Db	901	LWPTPSENDIVHLPRDMGHLQVDYRDLNRLHPSDESSLDSTASVVVPIIICLSIIIAFLFI	960
Qy	961	NOKKOWIPLCWTPTKPSLNNQLVSDCKKGTGRVQVDSOQRLRIAPDARFSGFYS	1020
Db	961	NOKKOWIPLCWTPTKPSLNNQLVSDCKKGTGRVQVDSOQRLRIAPDARFSGFYS	1020
Qy	1021	MQQNHLQADNFYQTV 1036	
Db	1021	MQQNHLQADNFYQTV 1036	
		AAU12242 standard; Protein; 1036 AA.	
		AAU12242	
		AC AAU12242;	
		AC AAU12242;	
		DT 24-OCT-2001 (first entry)	
		DE Human PRO4330 polypeptide sequence.	
		DE Human secretory and transmembrane; PRO; mammalian; cancer; lung;	
		KW breast; prostate; cervical; tumour necrosis factor-alpha; TNF-alpha;	
		KW cartilage; ear; proliferation; glucose; free fatty acid; skeletal muscle;	
		KW adipocyte; A-peptide; factor VIIa; gene therapy.	
		OS Homo sapiens.	
		XX WO200140466-A2.	
		PN 07-JUN-2001.	
		PD 01-DEC-2000; 2000WO-US32678.	
		XX 01-DEC-1999; 99WO-US28301.	
		PR	

XX AC AAU07141;
XX DT 24-OCT-2001 (first entry)
XX DE Human CRIM1 protein.
XX KW CRIM-1; Human; human chromosome 2p21-16.3; ophthalmological;
XX KW neuroprotective; renal; osteopathic; dental; vulnery; immunogen;
KW antibody; gene therapy; neurodegenerative disease; eye disorder;
KW cataract; bone morphogenic protein; BWG; renal disease; bone abnormality;
XX KW tooth abnormality; wound; S52.
OS Homo sapiens.
XX Key Location/Qualifiers
FH Peptide 1..17
FT /label= Signal_peptide
FT Domain 1..901
FT /label= Ectodomain
FT /note= "This sequence is specifically claimed in claim
FT 15"
FT Protein 18..1036
FT /label= Mature_CRIM1
FT Region 200..207
FT /note= "Conserved N-terminal motif"
FT Region 336..391
FT /label= CR_1
FT /note= "Cysteine rich repeat"
FT Region 403..456
FT /label= CR_2
FT /note= "Cysteine rich repeat"
FT Misc-difference 414
FT /note= "Encoded by GAC"
FT Region 608..662
FT /label= CR_3
FT /note= "Cysteine rich repeat"
FT Region 679..734
FT /label= CR_4
FT /note= "Cysteine rich repeat"
FT Region 753..808
FT /label= CR_5
FT /note= "Cysteine rich repeat"
FT Region 819..873
FT /label= CR_6
FT /note= "Cysteine rich repeat"
XX WO200138519-A1.
XX PD 31-MAY-2001.
XX 24-NOV-2000; 2000WO-AU01435.
XX 26-NOV-1999; 99AU-0004348.
XX (UYQU) UNIV QUEENSLAND.
XX Little M, Yamada T, Holmes G, Georgas K, Kolle G, Wilkinson L;
XX WPI: 2001-343951/36.
XX N-PSDB; AAS11601.
XX Nucleic acids from human chromosome 2p21-16.3 and the encoded peptide,
XX useful for preventing, diagnosing and treating e.g. eye disease,
XX especially cataract formation -
XX Claim 11; Fig 1; 169pp; English.
XX The invention relates to nucleic acids from human chromosome 2p21-16.3
XX and the encoded peptide (and mouse and chicken orthologues) that
XX comprises a PGCCPLP group, an insulin-like growth factor binding protein
XX (IGFBP)-like domain, cysteine-rich domains, an RGD (undefined) group
XX and a transmembrane domain. The protein, e.g. CRIM1, interacts with

CC peptides of the transforming growth factor superfamily. A composition
CC comprising an expression construct comprising the nucleic acids of the
CC invention or a mimetic which antagonises or mimics an activity of a CRIM1
CC polypeptide may be used in a method for modulating the biological
CC activity of a polypeptide of the bone morphogenic protein (BMP) family.
CC In this way they may be used to prevent or treat an eye disease,
CC especially cataract formation. they may also be used to treat
CC neurodegenerative diseases, renal and kidney disease, bone and tooth
CC abnormalities, wounds and skin damage, e.g. by use of the nucleic acid in
CC gene therapy by using antibodies directed against CRIM1 polypeptides.
CC The present sequence represents human CRIM1 (AKA S52).
XX
SQ Sequence 1036 AA;
Query Match 99.1%; Score 5901; DB 22; Length 1036;
Best Local Similarity 99.2%; Pred. No. 0;
Matches 1028; Conservative 3; Mismatches 5; Indels 0; Gaps 0;
Qy 1 MYLVAGDRGLAGCGHLLVSLGLLLPARSTRALVCLPDESKCEPRNRPGSIGVGC 60
Db 1 MYLVAGDRGLAGCGHLLVSLGLLLPARSTRALVCLPDESKCEPRNRPGSIGVGC 60
Qy 61 GCCYTASQGNESCGGTGIIYTCDRGLRCVIRPLNGDSLTYEAGVCEDENTDDQLL 120
Db 61 GCCYTASQGNESCGGTGIIYTCDRGLRCVIRPLNGDSLTYEAGVCEDENTDDQLL 120
Qy 121 GFPCNENLIAGCNITNGKCECNTIRTCNPFEPSPQDMCLSKRIEEKPCDSKARCE 180
Db 121 GFPCNENLIAGCNITNGKCECNTIRTCNPFEPSPQDMCLSKRIEEKPCDSKARCE 180
Qy 181 VQSPRPEDSVLIEGYAPGEGCPLPSRCVNCNPAGCLRVKQCPGNILVSKASGKPE 240
Db 181 VQSPRPEDSVLIEGYAPGEGCPLPSRCVNCNPAGCLRVKQCPGNILVSKASGKPE 240
Qy 241 CCDLYECKPVFVDCRTVECPVQGTACPPDSYETVRLTADGCTLTPECELSGLCGF 300
Db 241 CCDLYECKPVFVDCRTVECPVQGTACPPDSYETVRLTADGCTLTPECELSGLCGF 300
Qy 301 PVCEVGSTPRIVSRGDTGPKKCDVFECVNDTKPACVFNVEYDGMFMDNCRFCRCQ 360
Db 301 PVCEVGSTPRIVSRGDTGPKKCDVFECVNDTKPACVFNVEYDGMFMDNCRFCRCQ 360
Qy 361 GGVAICFTAQCGEINCERYVYVEGECPCVCEBDPVYPPNNPAGCYANGLIIAHGDRWREDD 420
Db 361 GGVAICFTAQCGEINCERYVYVEGECPCVCEBDPVYPPNNPAGCYANGLIIAHGDRWREDD 420
Qy 421 CTFQCQVNGERHCVATVCGQFTNPVKVPGCCPVCEPTIITVDPACGELSCTLTRK 480
Db 421 CTFQCQVNGERHCVATVCGQFTNPVKVPGCCPVCEPTIITVDPACGELSCTLTRK 480
Qy 481 DCINGFRDHNRCRTCCQINTQELCSERKQGTLCNCPFGELTDAQNCIECECRPPKKCR 540
Db 481 DCINGFRDHNRCRTCCQINTQELCSERKQGTLCNCPFGELTDAQNCIECECRPPKKCR 540
Qy 541 PIICDKYCPGLLLKNKHGCDICRCKKCPKPELSCKICPLGFGQDQSHGCLICKCRASASAG 600
Db 541 PIICDKYCPGLLLKNKHGCDICRCKKCPKPELSCKICPLGFGQDQSHGCLICKCRASASAG 600
Qy 601 PPIISGTCCLTVDGHGHHKNEESWHDGCRECYCLNGREMCALITCPVACGNPTTHPGQCCP 660
Db 601 PPIISGTCCLTVDGHGHHKNEESWHDGCRECYCLNGREMCALITCPVACGNPTTHPGQCCP 660
Qy 661 SCADDFVVKQPELSTPSICHAPGGGEYFVEGETWNIDSTCTCTCHSGRVLCTETECVPLL 720
Db 661 SCADDFVVKQPELSTPSICHAPGGGEYFVEGETWNIDSTCTCTCHSGRVLCTETECVPLL 720
Qy 721 QNPSRTQDSCPCQCTDQFPFRPSLRNNSVNPYCKNDEGDIPLAESWKPDCVCSICIDS 780
Db 721 QNPSRTQDSCPCQCTDQFPFRPSLRNNSVNPYCKNDEGDIPLAESWKPDCVCSICIDS 780
Qy 781 VISCFSESCPSVSCERPVLRKGQCCPYCIKDTIPKVVVCHFSKGAYADEERWLDSDCTHC 840
Db 781 VISCFSESCPSVSCERPVLRKGQCCPYCIKDTIPKVVVCHFSKGAYADEERWLDSDCTHC 840

QY 841 YCLOGTLCSTVSCPLPCVERP INVEGCCPCWCPMYVPEPTNPIPIEKTNRHGEVDLEVP 900
Db |||||
QY 841 YCLOGTLCSTVSCPLPCVERP INVEGCCPCWCPMYVPEPTNPIPIEKTNRHGEVDLEVP 900
Db |||||
QY 901 LMPPTSENDIVHLPRDMGHQVDYRDNRLHPSEDSLSLSDSIASVVVPIIICLSIIIAFLFI 960
Db |||||
QY 901 LMPPTSENDIVHLPRDMGHQVDYRDNRLHPSEDSLSLSDSIASVVVPIIICLSIIIAFLFI 960
Db |||||
QY 961 NQKKOWIPLLCWYRPTKPSLNNQLVSDCKKGTGRVQVDSQRMRLRTAEPDARFSGFYS 1020
Db |||||
QY 961 NQKKOWIPLLCWYRPTKPSLNNQLVSDCKKGTGRVQVDSQRMRLRTAEPDARFSGFYS 1020
Db |||||
QY 1021 MQQNHLQADNFYQTV 1036
Db |||||
QY 1021 MQQNHLQADNFYQTV 1036
Db |||||
RESULT 6
AAU07142
ID AAU07142 standard; Protein; 1037 AA.
XX AC AAU07142;
XX 24-OCT-2001 (first entry)
XX DE Mouse CRIM1 protein.
XX KW CRIM-1; Mouse; human chromosome 2p21-16.3; ophthalmological;
KW neuroprotective; renal; osteopathic; dental; vulnery; immunogen;
KW antibody; gene therapy; neurodegenerative disease; eye disorder;
KW cataract; bone morphogenic protein; BMG; renal disease; bone abnormality;
KW tooth abnormality; wound; S52.
XX OS Mus sp.
XX FH Key
FT Peptide 1...9
FT Protein /label= Signal_peptide
FT Protein 10..1037
FT Protein /label= Mature_CRIM1
FT Region 200..207
FT Region /note= "Conserved N-terminal motif"
FT Region 336..391
FT Region /label= CR_1
FT Region /note= "Cysteine rich repeat"
FT Region 403..456
FT Region /label= CR_2
FT Region /note= "Cysteine rich repeat"
FT Region 608..662
FT Region /label= CR_3
FT Region /note= "Cysteine rich repeat"
FT Region 679..734
FT Region /label= CR_4
FT Region /note= "Cysteine rich repeat"
FT Region 753..808
FT Region /label= CR_5
FT Region /note= "Cysteine rich repeat"
FT Region 819..873
FT Region /label= CR_6
FT Region /note= "Cysteine rich repeat"
XX WW0200138519-A1.
XX 31-MAY-2001.
XX 24-NOV-2000; 2000WO-AU01435.
XX 26-NOV-1999; 99AU-0004348.
XX (UYQU) UNIV QUEENSLAND.
XX Little M, Yamada T, Holmes G, Georgas K, Kolle G, Wilkinson L;

XX WPI: 2001-343951/36.
DR N-PSDB; AAS11602.
XX
FT Nucleic acids from human chromosome 2p21-16.3 and the encoded peptide,
PT useful for preventing, diagnosing and treating e.g. eye disease,
PT especially cataract formation -
XX
XX Claim 11; Fig 1; 169pp; English.
XX
CC The invention relates to nucleic acids from human chromosome 2p21-16.3
CC and the encoded peptide (and mouse and chicken orthologues) that
CC comprises a PGCCPLP group, an insulin-like growth factor binding protein
CC (IGFBP)-like domain, cysteine-rich domains, an RGD (undefined) group
CC and a transmembrane domain. The protein, e.g. CRIM1, interacts with
CC peptides of the transforming growth factor superfamily. A composition
CC comprising an expression construct comprising the nucleic acids of the
CC invention or a mimetic which antagonises or mimics an activity of a CRIM1
CC polypeptide may be used in a method for modulating the biological
CC activity of a polypeptide of the bone morphogenic protein (BMP) family.
CC In this way they may be used to prevent or treat an eye disease,
CC especially cataract formation. They may also be used to treat
CC neurodegenerative diseases, renal and kidney disease, bone and tooth
CC abnormalities, wounds and skin damage, e.g. by use of the nucleic acid in
CC gene therapy by using antibodies directed against CRIM1 polypeptides.
XX The present sequence represents mouse CRIM1 (AKA S52).
XX
SQ Sequence 1037 AA;

Query Match 90.8%; Score 5402.5; DB 22; Length 1037;
Best Local Similarity 88.5%; Pred. No. 6.7e-300;
Matches 918; Conservative 51; Mismatches 67; Indels 1; Gaps 1;

QY 1 MYLVAGDRLAGCGHLLVSLGLLLLPARSGTRALVCLPDCSEKCEEPNRFGSIVQVC 60
Db |||||
QY 1 MYLVAGDRLAGCGHLLVSLGLLLLPARSGTRALVCLPDCSEKCEEPNRFGSIVQVC 60
Db |||||
QY 61 GCCTCASQGNESCGTGGIYCTCDRLGRCVIRPPLNGSLTEYEGVCEDEWDQJLL 120
Db |||||
QY 61 GCCTCASQGNESCGTGGIYCTCDRLGRCVIRPPLNGSLTEYEGVCEDEWDQJLL 120
Db |||||
QY 121 GFKPCNENLIAGCNIINGKCECNIITRTCSNPFEPFSDQMCALSKALKRIEKEKPCSKARCE 180
Db |||||
QY 121 GFKPCNENLIAGCNIINGKCECNIITRTCSNPFEPFSDQMCALSKALKRIEKEKPCSKARCE 180
Db |||||
QY 181 VQSPRCPEDSVLIEGYAPPGGCCPLPSRCVNCNAGLCKRVCPQGNLNLVSKASKPGE 240
Db |||||
QY 181 VQSPRCPEDSVLIEGYAPPGGCCPLPSRCVNCNAGLCKRVCPQGNLNLVSKASKPGE 240
Db |||||
QY 241 CDDLYECKPVFGVDCRTVECPVQQTACPPDSYETQVRLTAGCCCTLPTRCECLSLGCGF 300
Db |||||
QY 241 CDDLYECKPVFGVDCRTVECPVQQTACPPDSYETQVRLTAGCCCTLPTRCECLSLGCGF 300
Db |||||
QY 301 PVCEVGSTPRIVSRGDTGPKGCDVPECVNDTKPACVFNNVEYDGDMPDMNCRCRCQ 360
Db |||||
QY 301 PVCEVGSTPRIVSRGDTGPKGCDVPECVNDTKPACVFNNVEYDGDMPDMNCRCRCQ 360
Db |||||
QY 361 GGVAICFTAQCGEINCERYYPVEGECPCVCDPVYFPNNPAGCYANGLILAHGDRWREDD 420
Db |||||
QY 361 GGVAICFTAQCGEINCERYYPVEGECPCVCDPVYFPNNPAGCYANGLILAHGDRWREDD 420
Db |||||
QY 421 CTFCCQVNGERHCVATVCGQTCTNPNKVPGECCPVEEPTIITVDPAGGELSNCTLTKK 480
Db |||||
QY 421 CTFCCQVNGERHCVATVCGQTCTNPNKVPGECCPVEEPTIITVDPAGGELSNCTLTKK 480
Db |||||
QY 481 DCINGFKRDHNGCORTCOQINTQELCSEKOGCTLNCPPGELTDAONCEICECRPKKCR 540
Db |||||
QY 481 DCINGFKRDHNGCORTCOQINTQELCSEKOGCTLNCPPGELTDAONCEICECRPKKCR 540
Db |||||
QY 541 PIICDKYCPGLGLKNKHGCDICRCKKCPSELCKSCIPCLGFGQDSHGCLICKREASAG 600
Db |||||
QY 541 PIICDKYCPGLGLKNKHGCDICRCKKCPSELCKSCIPCLGFGQDSHGCLICKREASAG 600
Db |||||

QY 601 PPLSTCLTVDGHHKHEESWHDGCRECYCLNGREMCALITCPVPACGNPTIHPGQCCP 660
Db 601 PVLSTCLTVDGHHKHEESWHDGCRECYCLNGREMCALITCPVPACGNPTIHPGQCCP 660
QY 661 SCADDFVQKPELSTPSICHAPGGEYFVEGETWINDSCTQCHSGRVLCTEVCPPLLC 720
Db 661 SCTDDFVQKPELSTPSICHAPGGEYFVEGETWINDSCTQCHSGRVLCTEVCPPLLC 720
QY 721 QNPSRTQDSCCPQCTDQPFRLSRNSVPNYCKNDEGDFILAAESWKPVDVCTSGICIDS 780
Db 721 QNPSRTQDSCCPQCTDQPFRLSRNSVPNYCKNDEGDFILAAESWKPVDVCTSGICIDS 780
QY 781 VISCFSSECPVSCRPVLRKQCCPYCIKDTIPKVVCHFSKAYADEERWDLDSCTHC 840
Db 781 AISCYSECPVACERPLVRKQCCPYCLEDTIPKVVCHFSKAYADEERWDLDSCTHC 840
QY 841 YCLOGQTLCTVSCPLPCVPEPINVEGSCPCMPYVPEPINPIEKTNNHGEVDLEVP 900
Db 841 YCLOGQTLCTVSCPLPCVPEPINVEGSCPCMPYVPEPINPIEKTNNHGEVDLEVP 900
QY 901 LWPTSENDIVHLPDMGHQVDYRD-NRLHPSSESSLDSTIASVVVPIIICLSIIIAFLF 959
Db 901 MWPTSENDIHLPRDMGHQVDYRDNNRLHPSSESSLDSTIASVVVPIIICLSIIIAFLF 960
QY 960 INOKKQWIPLLCWYRTPKPSLNQVSDCKKGTFRVQVDSQRMRLTAEPDARFSGFY 1019
Db 961 INOKKQWIPLLCWYRTPKPSLNQVSDCKKGTFRVQVDSQRMRLTAEPDARFSGFY 1020
QY 1020 SMOKQNHQADNFYQTV 1036
Db 1021 SMOKQNHQADNFYQTV 1037

RESULT 7
AAU07143
ID AAU07143 standard; Protein; 1048 AA.
XX AAU07143;
AC AAU07143;
XX 24-OCT-2001 (first entry)
XX Chicken CRIM1 protein.
DE CRIM-1; Chicken; human chromosome 2p21-16.3; ophthalmological;
XX neuroprotective; renal; osteopathic; dental; vulnary; immunogen;
KW antibody; gene therapy; neurodegenerative disease; eye disorder;
KW cataract; bone morphogenic protein; BMG; renal disease; bone abnormality;
KW tooth abnormality; wound; S52.
XX Gallus gallus.
OS

Key Location/Qualifiers
FH 212..219
FT Region /note= "Conserved N-terminal motif"
FT Region 348..402
FT /label= CR_1
FT /note= "Cysteine rich repeat"
FT Region 415..468
FT /label= CR_2
FT /note= "Cysteine rich repeat"
FT Region 620..674
FT /label= CR_3
FT /note= "Cysteine rich repeat"
FT Region 691..746
FT /label= CR_4
FT /note= "Cysteine rich repeat"
FT Region 765..820
FT /label= CR_5
FT /note= "Cysteine rich repeat"
FT Region 831..885
FT /label= CR_6
FT /note= "Cysteine rich repeat"

FN WO200138519-A1.
XX 31-MAY-2001.
XX 24-NOV-2000; 2000WO-AU01435.
XX 26-NOV-1999; 99AU-0004348.
XX (UYQU) UNIV QUEENSLAND.
XX Little M, Yamada T, Holmes G, Georgas K, Kolle G, Wilkinson L;
PI WPI; 2001-343951/36.
XX N-PSDB; AAS11603.
XX Nucleic acids from human chromosome 2p21-16.3 and the encoded peptide,
PT useful for preventing, diagnosing and treating e.g. eye disease,
FT especially cataract formation -
XX Claim 11; Fig 1; 169pp; English.
XX The invention relates to nucleic acids from human chromosome 2p21-16.3
CC and the encoded peptide (and mouse and chicken orthologues) that
CC comprises a PGECCP group, an insulin-like growth factor binding protein
CC (IGFBP)-like domain, cysteine-rich domains, an RGD (undefined) group
CC and a transmembrane domain. The protein, e.g. CRIM1, interacts with
CC peptides of the transforming growth factor superfamily. A composition
CC comprising an expression construct comprising the nucleic acids of the
CC invention or a mimetic which antagonises or mimics an activity of a CRIM1
CC polypeptide may be used in a method for modulating the biological
CC activity of a polypeptide of the bone morphogenic protein (BMP) family.
CC In this way they may be used to prevent or treat an eye disease,
CC especially cataract formation. They may also be used to treat
CC neurodegenerative diseases, renal and kidney disease, bone and tooth
CC abnormalities, wounds and skin damage, e.g. by use of the nucleic acid in
CC gene therapy by using antibodies directed against CRIM1 polypeptides.
CC The present sequence represents chicken CRIM1 (AKA S52).
XX

Query Match 83.5%; Score 4969; DB 22; Length 1048;
Best Local Similarity 80.6%; Pred. No. 3.4e-275;
Matches 846; Conservative 80; Mismatches 109; Indels 14; Gaps 5;
Qy 1 MYLVA-----GDRGLACGG-HLLVVS--LLGILLPLPARSGTRALVCLPCDESKCEEP 48
Db 1 MYLAASAGRPRPGDGGGGGWHLAAAGWLLLLALLGQPGTRALVCLPCDESKCEEP 60
Qy 49 RNRPGSIVQGVCGCYTCASOGNESCGETFGIYCTCDRGLRCVTRPPLNGDSLTEYAGV 108
Db 61 KSCPGIIVLIGCGCCFMCARQNRNESCQGVYGLHACDRGLRCVTRPPLNGDSLTEYAGV 120
Qy 109 CEDENWTDQLLGFKPCNENIAGCNITNGKCEINTTCNPFEPFSDQMCLSAKRIE 168
Db 121 CEDENWDDQLLGFEPNENITGNTIINGKCDCTTTCNPFEPFSDTCLSAKRIE 180
Qy 169 BEKPDCKARCEVOFSPRCPEDSVLIEGYAPPGECCPLPSRCVNCNAGCLRKVCQPNLN 228
Db 181 BEKPDCKARCEVOFSPRCPEDSVLIEGYAPPGECCPLPSRCVNCNAGCLRKVCQPNLN 240
Qy 229 ILVSKASGKPGCCDLYECKPVFGVDCRTVECPVQQTACPPDSYETQVRLTAGCCCTLP 288
Db 241 ILVSKASGKPGCCDLYECKPVFVSDCTVECPVQVQVWCPPLDSYETQVRLTAGCCCTLP 300
Qy 289 TRCECLSLGCGFPVCEVGSTPRIVSRGDTGPKGCCDFECVNDTKPACVFNNVYYDGM 348
Db 301 TRCECLSLGCGFPVCEAGSVQIVSRGDTGPKGCCDFECVNEVKPTCFNSMEYDGM 360
Qy 349 FRMDNCRFCRCGGVAICFTAQCGEINCERYYPVEGECPCVCEDPVYFPNNPACGYANGL 408
Db 361 FRMDACRFRCCGGVSICTFSAQCGELHCDYVPEGECPCVCEDPVYFPNNPACGYANG 420
Qy 409 ILAHGDRWRDDCTFCQCVNGERHCVATVCGTCTNPNKVPGECCPCVCEEPTIITVPPA 468

Db 421 IQAHDRWREDDCTFCQINGNPHCVATACGSCNLPVKVPGECPCVCEETIYIGPPT 480
QY 469 CGELSNCTLTTRKDCINGFKRKHNGCTQCINTQELCSERKQGTFLNCPFGFLDAQNCE 528
Db 481 CELLVNCTLTEDKCIYSFKLDONGRCICQCKTRELCITGLISGCLDSCSFGQDAHNC 540
QY 529 ICECRPRPKKCRPIIDKCYPLGLLKNKHGCDICRCKKCPSELCSKICPLGFOODSHGCL 588
Db 541 IQCRRPRPKKCKPIVCDKCYCPFGYLKKNKHGCEICRCKKCPMPCKGKICPMGFQNSHGCV 600
QY 589 ICKCREASASAGPPTLSGTCLTVDGHGHHKNEESWHDGRCYCYNLNGREMCALITCPVPAC 648
Db 601 ICKCREATASLMPVYKTSGLSMDGRHNEESWHDGRCYCYNLNGREMCALITCPVPNC 660
QY 649 GNPTTHPGQCCPCADDVWQKPELSTPSICHAPGEVFEVGETWNIIDSCCTQCTCHSGRV 708
Db 661 GNPTTHPGQCCPCDEIIVQKPELSTPSICHAPGEVFEVGETWNIIDSCCTQCTCHSGRV 720
QY 709 LCETEVCPPLLCQNPSTRQDSCCPQCTDQFPRPSLSRNSVNPYCKNDEGDIFLAASWK 768
Db 721 LCETEVCPPLLCQNPSTRQDSCCPQDEPLQPSLSSNVSMPSYCKNDEGDIFLAASWK 780
QY 769 PVCTSCICIDSVISCFSESPSCERPVLKRGQCCPYCIKDTIPKKVCHFGSKAYAD 828
Db 781 PVNCTSCICMDGVIRCYSESPVSCERPVLKRGQCCPYCIETDTPKKVCHFGKTYAD 840
QY 829 ERWDLDSCTHCYCLQGTCLSTVSCPLPCVPERINVEGSCPCMPENYVPEPTNPIEK 888
Db 841 ERWDLDSCTHCYCLQGTCLSTVSCPLPCVPERINVEGSCPCMPENYVPEPTNPIEK 900
QY 889 TNHRGEVLEVPWLPPTSENDIVHLPRDMGHQVDYRD-NRLHPSEDSLSIASVVVPI 947
Db 901 TNHRGDVELEVPWLPPTSENDIHLHRDMNHLQGEYRSCNGPHPSADASVSSVALVTPI 960
QY 948 ICLSIITAFINOKKOWIPLLCWRYPTTPSSLNOLVSVDCCKGTRVQVDSQRMRLR 1007
Db 961 TIALVIVFLINOKKOWIPVSC-YKAPTTPKPSCLNQLVYVDCCKGTRVQVDSQRMRLR 1019
QY 1008 IAEPAFSGFYSMQKQNLQADNEYQTV 1036
Db 1020 IADPDSRYSGFYSMQKQNLQADNEYQTV 1048

RESULT 8
AAB61140
ID AAB61140 standard; Protein; 732 AA.
XX AC AAB61140;
XX DT 30-MAR-2001 (first entry)
XX DE Human NOV10 protein.
XX KW Human; NOVX; antiinflammatory; cytostatic; neuroprotective;
KW cerebroprotective; immunomodulator; vulnerary; vasotropic; gene therapy;
KW hyperplasia; tumour; restenosis; psoriasis; Dupuytren's contracture;
KW diabetes; rheumatoid arthritis; cerebral oedema; Alzheimer's disease.
XX OS Homo sapiens.
XX PN WO2000075321-A2.
XX PD 14-DEC-2000.
XX PF 01-JUN-2000; 2000WO-US15303.
XX PR 03-JUN-1999; 99US-0137322.
XX PR 16-MAR-2000; 2000US-0189810.
XX PR 22-MAR-2000; 2000US-0191158.
XX PR 30-MAR-2000; 2000US-0193086.
XX PR 31-MAY-2000; 2000US-0137322.
XX

(CURA-) CURAGEN CORP.
Shimkets RA, Fernandes E, Herrman J, Vernet C;
WPI; 2001-102403/11.
N-PSDB; AAF27858.

New NOVX polypeptides and polynucleotides, useful in gene therapy, as a diagnostic marker, protein therapeutic, antibody or small molecule drug target for treating immune, proliferative and metabolic diseases and wound healing
Claim 1; Page 39-42; 194pp; English.

The present sequence is a new isolated polypeptide (NOVX). The NOVX polypeptides, NOVX nucleic acids, and anti-NOVX antibodies are useful for treating or preventing NOVX-associated disorders. They are also useful for determining the presence of or a predisposition to a disease associated with altered levels of the NOVX polypeptide or nucleic acid. These NOVX-associated disorders include hyperplasias, tumours, restenosis, psoriasis, Dupuytren's contracture, diabetic complications, rheumatoid arthritis, cerebral lesions, diabetic neuropathies, cerebral oedema, senile dementia or Alzheimer's disease. The NOVX polynucleotides are especially useful in gene therapy. Specifically, NOVX is useful as a diagnostic marker or prognostic marker, protein therapeutic and antibody target or small molecule drug target to treat disorders in the immune response pathway, thyroid and metabolic diseases, bone metabolic disorders, diseases of the pancreas (e.g. diabetes or digestive disorders), proliferative diseases, or tissue regeneration and development (e.g. wound healing or treatment of burns).
XX Sequence 732 AA;
QY Query Match 70.0%; Score 4167; DB 22; Length 732;
Best Local Similarity 98.6%; Pred. No. 1.2e-229;
Matches 711; Conservative 3; Mismatches 7; Indels 0; Gaps 0;

QY 159 MCLSALKRIEERKDCSKARCEVQFSPRCPEDSVLIEGYAPPGCECCPLPSRCVNCNAGCL 218
Db 1 MCLSALKRIEERKDCSKARCEVQFSPRCPEDSVLIEGYAPPGCECCPLPSRCVNCNAGCL 60

QY 219 RKVCQPGNLLILVSKASGKPGCECCDLYECKPVFGVDCRTVECPVQQTACPPDSYETQVR 278
Db 61 RKVCQPGNLLILVSKASGKPGCECCDLYECKPVFGVDCRTVECPVQQTACPPDSYETQVR 120

QY 279 LTADGGCTLTTRCECLSGLCGFPVCEVSGTSPRIVSRGDTGTPGCCDVFECVNDTKPACVF 338
Db 121 LTADGGCTLTTRCECLSGLCGFPVCEVSGTSPRIVSRGDTGTPGCCDVFECVNDTKPACVF 180

QY 339 NNVEYDGDMDFRMDCNRCFCRCQGVVAICFTAQCGEINCERYVYVPEGECPCVCDPVYFPN 398
Db 181 NNVEYDGDMDFRMDCNRCFCRCQGVVAICFTAQCGEINCERYVYVPEGECPCVCDPVYFPN 240

QY 399 NPAGCYANGLILAHGDRWRDEDDCTFCQVNGERHCVATVCGQTCTNPVKVPGECPCVCEE 458
Db 241 NPAGCYANGLILAHGDRWRDEDDCTFCQVNGERHCVATVCGQTCTNPVKVPGECPCVCEE 300

QY 459 PTITVDPACGELSNTLTTRKDCINGFKRDHNGCTCQCINTQELCSERKQGTCLNCPF 518
Db 301 PTITVDPACGELSNTLTTRKDCINGFKRDHNGCTCQCINTQELCSERKQGTCLNCPF 360

QY 519 GFLTDAQNCEICRPRPKCRPIICDKYCPGLLKNKHGCDICRCKKPELSCSKICPL 578
Db 361 GFLTDAQNCEICRPRPKCRPIICDKYCPGLLKNKHGCDICRCKKPELSCSKICPL 420

QY 579 GFQODSHGCLICKREASASAGPPIILSGTCLTVDGHGHHKNEESWHDGRCYCYNLNGREMC 638
Db 421 GFQODSHGCLICKREASASAGPPIILSGTCLTVDGHGHHKNEESWHDGRCYCYNLNGREMC 480

QY 639 ALITCPVPACGNPTIHPGCCPCSCADDVWQKPELSTPSICHAPGEVFEVGETWNIIDSC 698
Db 481 ALITCPVPACGNPTIHPGCCPCSCADDVWQKPELSTPSICHAPGEVFEVGETWNIIDSC 540

QY 699 TQCTCHSGRVLCTETVCPPLLQCNPSRTQDSCCPQCTDQPPRPSLSRNNNSVNPYCKNDEG 758
Db 541 TQCTCHSGRVLCTETVCPPLLQCNPSRTQDSCCPQCTDQPPRPSLSRNNNSVNPYCKNDEG 600
QY 759 DIFLAESKPKDVCTSCICIDSVISCFSSCPVSRKGVKGGCCPCYCIKDTTPKKV 818
Db 601 DIFLAESKPKDVCTSCICIDSVISCFSSCPVSRKGVKGGCCPCYCIKDTTPKKV 660
QY 819 CFFSGKAYADEERWDLDSCTHYCQLQGQTLCTVSCPPPLPCVPEINVEGSCCPMCPMKV 878
Db 661 CFFSGKAYADEERWDLDSCTHYCQLQGQTLCTVSCPPPLPCVPEINVEGSCCPMCPVSP 720
QY 879 P 879
Db 721 P 721
RESULT 9
ABG66681
ID ABG66681 standard; Protein; 503 AA.
XX
AC ABG66681;
XX
DT 30-AUG-2002 (first entry)
XX
DE Human novel polypeptide #16.
XX
KW Human; inflammatory condition; shock; sepsis; immune response;
KW cancer; wound healing; central nervous system disease; haematopoiesis;
KW peripheral nervous system disease; amyotrophic lateral sclerosis; tendon;
KW myeloid cell disorder; lymphoid cell disorder; platelet disorder; bone;
KW cartilage; ligament; nerve tissue; ulcer; osteoporosis; osteoarthritis;
KW bone degenerative disorder; periodontal disease; reperfusion injury;
KW lung fibrosis; liver fibrosis; autoimmune disorder; bacterial infection;
KW allergic condition; thrombolysis; thrombosis; coagulation disorder;
KW fungal infection.
XX
OS Homo sapiens.
XX
PN WO200244340-A2.
XX
PD 06-JUN-2002.
XX
PF 30-NOV-2001; 2001WO-US47004.
XX
PR 30-NOV-2000; 2000US-0028952.
XX
PA (HYSE-) HYSEQ INC.
XX
PI Tang YT, Goodrich RW, Liu C, Zhou P, Asundi V, Wang J, Wang D;
PI Yamazaki V, Ujwal ML, Drmanac RT;
XX
DR WPI: 2002-508509/54.
DR N-PSDB; ABK94905.
XX
PT Novel nucleic acids and polypeptides for diagnosis, treatment of
PT inflammatory, autoimmune, nervous system, myeloid or lymphoid cell
PT disorders, cancer and promoting wound healing -
XX
PS Claim 10; Page 580-581; 672pp; English.
XX
CC The invention relates to human novel polynucleotides and associated
CC polypeptides. The polynucleotides and polypeptides are useful for
CC treating inflammatory conditions such as arthritis, nephritis, Crohn's
CC disease, ischaemia-reperfusion injury, shock, sepsis, immune responses
CC and cancer and for promoting wound healing. The sequences are used to
CC induce the proliferation of neural cells and regeneration of nerve and
CC brain tissue, and are useful for the treatment of central and peripheral
CC nervous system diseases and neuropathies, such as Alzheimer's disease,
CC Parkinson's disease, Huntington's disease and amyotrophic lateral
CC sclerosis. The sequences are involved in chemotactic or chemokinetic
CC activity, regulation of haematopoiesis, treatment of myeloid or lymphoid
CC cell disorders and platelet disorders such as thrombocytopenia,

CC regeneration of bone, cartilage, tendon, ligament and/or nerve tissue
CC growth, tissue repair, healing of burns, incisions, ulcers, treatment of
CC osteoporosis, osteoarthritis, bone degenerative disorders and periodontal
CC disease. The sequences of the invention are also useful for gut
CC protection or regeneration and treatment of lung or liver fibrosis,
CC reperfusion injury in various tissues, immune deficiencies and disorders
CC including severe combined immunodeficiency (SCID), bacterial or fungal
CC infections, autoimmune disorders e.g. multiple sclerosis and myasthenia
CC gravis, allergic conditions such as asthma, thrombolysis or thrombosis
CC and coagulation disorders. Sequences ABG66666-ABG66758 represent human
CC novel polypeptides of the invention.
XX
SQ Sequence 503 AA;
Query Match 47.5%; Score 2828.5; DB 23; Length 503;
Best Local Similarity 98.0%; Pred. No. 1.5e-153;
Matches 493; Conservative 2; Mismatches 7; Indels 1; Gaps 1;
QY 1 MYLVAGDRGLAGCGHLLVSLGLLLPARSGTRALVCLPCDESKCEPRNRPISVQGV 60
Db 1 MYLVAGDRGLAGCGHLLVSLGLLLPARSGTRALVCLPCDESKCEPRNRPISVQGV 60
QY 61 GCCYTCAQSNESCGGTGFIYGTCDRLGRCVIRPPLNGDSLTYEAGVCEDEWTDQLL 120
Db 61 GCCYTCAQSNESCGGTGFIYGTCDRLGRCVIRPPLNGDSLTYEAGVCEDEWTDQLL 120
QY 121 GFKPCNENLAGCNITNGKCECNTIRTCNPFPPSODMCLSAKRIEEKPDCSKARCE 180
Db 121 GFKPCNENLAGCNITNGKCECNTIRTCNPFPPSODMCLSAKRIEEKPDCSKARCE 180
QY 181 VQSPRCPEDSVLTIEGYAPGECCLPSPRCVNCNPAGCLRKVCQPNLNILVSKASKGPE 240
Db 181 VQSPRCPEDSVLTIEGYAPGECCLPSPRCVNCNPAGCLRKVCQPNLNILVSKASKGPE 240
QY 241 CCDIYECKPFGVDCRVCEPTVQQTARCPDDSYETQVRLTADGCCTLPTRCECLSLG 299
Db 241 CCDIYECKPFGVDCRVCEPTVQQTARCPDDSYETQVRLTADGCCTLPTRCECLSLG 300
QY 300 FPCVEGSTPRIVSRGDTGKCDVFECVNDTKPACVFNNVYDGMFMDNCFRC 359
Db 301 FPCVEGSTPRIVSRGDTGKCDVFECVNDTKPACVFNNVYDGMFMDNCFRC 360
QY 360 QGGVAICFTAQCGEINCERYVYVPEGCCPCVCEDPVYFPNPNAGCYANGLILAHGDRW 419
Db 361 QGGVAICFTAQCGEINCERYVYVPEGCCPCVCEDPVYFPNPNAGCYANGLILAHGDRW 420
QY 420 DCTFCQCVNGERHCVATVCGTCTNPNVKVPGCECPVCEPTIITVDPAGGELSNTLTR 479
Db 421 DCTFCQCVNGERHCVATVCGTCTNPNVKVPGCECPVCEPTIITVDPAGGELSNTLTR 480
QY 480 KDCINGFKRDHNGCRTCCQINTQ 502
Db 481 KDCINGFKRDHNGCRTCCQINSE 503
RESULT 10
AAAY82775
ID AAAY82775 standard; Protein; 400 AA.
XX
AC AAAY82775;
XX
DT 19-JUN-2000 (first entry)
XX
DE Human chordin related protein (Clone dj167_2).
XX
KW Chordin related protein; cartilage; bone; connective tissue;
KW periodontal disease; osteoporosis; burn; incision; ulcer; neuron;
KW astrocyte; glial cell; transplantation; nerve; epidermis; muscle;
KW liver; brain; lung; cardiac; pancreas; kidney; growth;
KW differentiation; TGF-Beta; angiogenesis; chemotaxis;
KW chemoattraction; collagen synthesis; fibrosis; cell adhesion;
KW cell migration; fertility; reproduction; haematopoiesis;
KW erythroid cell; tumour; dietary supplement; growth medium.

CC isolated from adult placenta, adult retina, foetal brain, foetal kidney,
CC adult blood, adult brain, adult thyroid, adult bladder, adult neural
CC tissue, adult testes, and adult lymph node cDNA libraries. The human
CC secreted proteins, and the polynucleotides encoding them, are predicted
CC to have biological activities which would make them suitable for
CC treating, preventing or ameliorating medical conditions in humans and
CC animals. Suggested activities include nutritional activity, cytokine
CC and cell proliferation/differentiation activity, immune stimulating
CC (e.g. as vaccines) or suppressing activity, haematopoiesis regulating
CC activity, tissue growth activity, activin/inhibin activity,
CC chemotactic/chemokinetic activity, haemostatic and thrombolytic
CC activity, receptor/ligand activity, anti-inflammatory activity,
CC cadherin/tumour invasion suppressor activity, and tumour inhibition
CC activity. The polynucleotides are also stated to be useful for gene
CC therapy. AAZ33316 to AAZ33373 encode human secreted proteins, and
CC AAY52998 to AAY53060 represent human secreted proteins, given in the
CC present invention.
XX
SQ Sequence 400 AA;

Query Match 37.9%; Score 2254; DB 21; Length 400;
Best Local Similarity 99.8%; Pred. No. 6.6e-121;
Matches 399; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 637 MCALITCPVACGNPTIHPGCCPCSCADDFVVKPELSTPSICHAPGGEYFVEGETWNID 696
DB 1 MCALITCPVACGNPTIHPGCCPCSCADDFVVKPELSTPSICHAPGGEYFVEGETWNID 60
QY 697 SCTQCTCHSGRVLCEVECPPLLQCNPSRTQDSCCPQCTDQFRLSLRNSVNPYCKND 756
DB 61 SCTQCTCHSGRVLCEVECPPLLQCNPSRTQDSCCPQCTDQFRLSLRNSVNPYCKND 120
QY 757 EGDIFLAESWPKPDVCTCICIDSVISCFSESCPSVSCRPVLRGQCCPYCIKDTIPKK 816
DB 121 EGDIFLAESWPKPDVCTCICIDSVISCFSESCPSVSCRPVLRGQCCPYCIEDTIPKK 180
QY 817 VVCHFSGKAYADERWDLDSCTHCYCLQQTLCSTVSCPPFCVPIVNEGSCCPMCPBM 876
DB 181 VVCHFSGKAYADERWDLDSCTHCYCLQQTLCSTVSCPPFCVPIVNEGSCCPMCPBM 240
QY 877 VYPEPTNIPTEKTHRGVEDLVEPLMPTSENDIVHLPRDMGHQVDVDRNLRHSEDS 936
DB 241 VYPEPTNIPTEKTHRGVEDLVEPLMPTSENDIVHLPRDMGHQVDVDRNLRHSEDS 300
QY 937 LDSIASVWVPIIICLSIIIAFLINQKKQWIPLLCWYRTPTKPSLNQLVSDVCKKQTR 996
DB 301 LDSIASVWVPIIICLSIIIAFLINQKKQWIPLLCWYRTPTKPSLNQLVSDVCKKQTR 360
QY 997 VQVDSQRMRLTAEPDARFSGFYSMQKNHLQADNFYQTV 1036
DB 361 VQVDSQRMRLTAEPDARFSGFYSMQKNHLQADNFYQTV 400

RESULT 12
AAB40954
ID AAB40954 standard; Protein; 322 AA.
XX
AC AAB40954;
XX
DT 08-FEB-2001 (first entry)
XX
DE Human ORFX ORF718 polypeptide sequence SEQ ID NO:1436.
XX
KW Human; open reading frame; ORFX; detection; cytostatic; hepatotropic;
KW vulnery; antiporiatic; antiparkinsonian; nootropic; neuroprotective;
KW anticonvulsant; osteopathic; antiarthritic; immunosuppressant; cardiant;
KW immunostimulant; thrombolytic; coagulant; vasotropic; antidiabetic;
KW hypotensive; dermatological; immunosuppressive; antiinflammatory;
KW antiviral; antibacterial; antifungal; antineumatic; antithyroid;
KW antianaemic; gene therapy; cancer; proliferative disorder; hypertension;
KW neurodegenerative disorder; osteoarthritis; graft vs host disease;
KW cardiovascular disease; diabetes mellitus; hypothyroidism; SCID; AIDS;
KW cholesterol ester storage; systemic lupus erythematosus; infection;

severe combined immunodeficiency; malaria; autoimmune disorder; asthma;
allergy; aplastic anaemia; nocturnal haemoglobinuria; burn; wound;
bone damage; cartilage damage; antiinflammatory disease; coagulation;
thrombosis; contraceptive.
OS Homo sapiens.
XX WO200058473-A2.
XX 05-OCT-2000.
XX 31-MAR-2000; 2000WO-US086621.
XX 31-MAR-1999; 99US-0127607.
PR 02-APR-1999; 99US-0127636.
PR 05-APR-1999; 99US-0127728.
PR 30-MAR-2000; 2000US-0540763.
XX
PA (CURA-) CURAGEN CORP.
XX
XX Shimkets RA, Leach M;
XX WPI: 2000-602362/57.
DR N-PSDB; AAC75163.
DR
XX Novel nucleic acids and peptides derived from open reading frame X,
PT useful for treating e.g. cancers, proliferative disorders,
PT neurodegenerative disorders and cardiovascular disease -
XX
PS Claim 11; Page 1214-1215; 5507pp; English.
XX
CC AAC74446 to AAC77606 encode the proteins given in AAB40237 to AAB43397,
CC which represent the human ORFX open reading frames 1 to 3161. The ORFX
CC sequences have activities such as: cytostatic; hepatotropic; vulnery;
CC antiporiatic; antiparkinsonian; nootropic; neuroprotective;
CC osteopathic; anticonvulsant; antiarthritic; immunosuppressant;
CC immunostimulant; cardiant; thrombolytic; coagulant; vasotropic;
CC antidiabetic; hypotensive; dermatological; immunosuppressive;
CC antinflammatory; antibacterial; antiviral; antifungal; antineumatic;
CC antithyroid; and antianaemic. The sequences can be used for determining
CC the presence of or predisposition to, or preventing or treating
CC pathological conditions associated with an ORFX-associated disorder. The
CC nucleic acids can be used to express ORFX proteins in gene therapy.
CC vectors. The proteins and nucleic acids may be used to treat cancers,
CC proliferative disorders, neurodegenerative disorders, osteoarthritis,
CC graft vs host disease, cardiovascular disease, diabetes mellitus,
CC hypertension, hypothyroidism, cholesterol ester storage, systemic lupus
CC erythematosus, severe combined immunodeficiency (SCID), AIDS, viral,
CC bacterial or fungal infection, malaria, autoimmune disorders, asthma,
CC allergies, aplastic anaemia, burns, wounds, bone and cartilage damage,
CC nocturnal haemoglobinuria, antiinflammatory disease; to enhance
CC coagulation; to inhibit thrombosis; and as a contraceptive.
XX
SQ Sequence 322 AA;

Query Match 26.5%; Score 1575; DB 21; Length 322;
Best Local Similarity 83.0%; Pred. No. 2.6e-82;
Matches 279; Conservative 3; Mismatches 28; Indels 26; Gaps 4;

QY 295 SGLCGFPVCEVGSTPRIVSRGDTGPKCCDVFECVNDTKPACVFNNVVEYDGMFRMDNC 354
DB 2 SGLCGFPVCEVGSTPRIVSRGDTGPKCCDVFECVNDTKPACVFNNVVEYDGMFRMDNC 61
QY 355 RFRCCQGVAICTTAQCGEINCERYVYVEGECPCVCEDDPVYFPNPNAGCYANGLILAHGD 414
DB 62 RFRCCQGVAICTTAQCGEINCERYVYVEGECPCVCEDDPVYFPNPNAGCYANGLILAHGD 120
QY 415 RWREDDCTFCOCVNGERHCVATVCGQCTNPVKVPVGGCCVCEPPTIITVDPACGELSN 474
DB 121 RWREDDCTFCOCVNGERHCVATVCGQCTNPVKVPVGGCCVCEPPTIITVDPACGELSN 180
QY 475 CTLTRKOCINGFKRHNGCRFTCCQINTQELCSERKQCTLNCPCFLTDAQNCIEICRCP 534
DB 475 CTLTRKOCINGFKRHNGCRFTCCQINTQELCSERKQCTLNCPCFLTDAQNCIEICRCP 534

Db 181 CTLTKDCINGPKRDHNGCRTCQCINTBELGASERKQGCTLNCPPFGFLTDAQNCEICECRP 240

QY 535 RPKKCRPIICDKYCPGLGLLNKHGCDICRCKKCPPELSCKICPLGFOODSHGCLICKCRE 594

Db 241 RPKKCRPIICDKYCPGLGLLNKHGCDICRCKKCPPELSCKICPLGFOODSHGCLICKCRE 286

QY 595 ASASAGPPILS-----GTLCTVDGH--HHKNEESW 622

Db 287 ---PAGQSRLSYLQVQRGGLCFSWATHPVGHLSHRGW 319

RESULT 13

AAU07149

ID AAU07149 standard; Protein; 872 AA.

XX

AC AAU07149;

XX

DT 24-OCT-2001 (first entry)

XX

DE C. elegans CRIM1 protein.

XX

KW CRIM-1; human chromosome 2p21-16.3; ophthalmological;

KW neuroprotective; renal; osteopathic; dental; vulnery; immunogen;

KW antibody; gene therapy; neurodegenerative disease; eye disorder;

KW cataract; bone morphogenic protein; BMG; renal disease; bone abnormality;

KW tooth abnormality; wound; S52.

XX

OS Caenorhabditis elegans.

XX

Key Location/Qualifiers

FT Region 31..38

FT /note= "Conserved N-terminal motif"

FT Region 172..228

FT /label= CR_1

FT /note= "Cysteine rich repeat"

FT Region 231..285

FT /label= CR_2

FT /note= "Cysteine rich repeat"

FT Region 424..469

FT /label= CR_3

FT /note= "Cysteine rich repeat"

FT Region 509..566

FT /label= CR_4

FT /note= "Cysteine rich repeat"

FT Region 584..635

FT /label= CR_5

FT /note= "Cysteine rich repeat"

FT Region 645..698

FT /label= CR_6

FT /note= "Cysteine rich repeat"

XX

WO2001138519-A1.

XX

31-MAY-2001.

XX

24-NOV-2000; 2000WO-AU01435.

XX

26-NOV-1999; 99AU-0004348.

XX

(UYQU) UNIV QUEENSLAND.

XX

Little M, Yamada T, Holmes G, Georgas K, Kolle G, Wilkinson L;

PI WPI; 2001-343951/36.

XX

Nucleic acids from human chromosome 2p21-16.3 and the encoded peptide,

PT useful for preventing, diagnosing and treating e.g. eye disease,

PT especially cataract formation -

XX

Example 14; Fig 4; 169pp; English.

PS

XX

The invention relates to nucleic acids from human chromosome 2p21-16.3

CC

and the encoded peptide (and mouse and chicken orthologues) that

CC comprises a pGECPLP group, an insulin-like growth factor binding protein

CC (IGFBP)-like domain, cysteine-rich domains, an RGD (undefined) group

CC and a transmembrane domain. The protein, e.g. CRIM1, interacts with

CC peptides of the transforming growth factor superfamily. A composition

CC comprising an expression construct comprising the nucleic acids of the

CC invention or a mimetic which antagonises or mimics an activity of a CRIM1

CC polypeptide may be used in a method for modulating the biological

CC activity of a polypeptide of the bone morphogenic protein (BMP) family.

CC In this way they may be used to prevent or treat an eye disease,

CC especially cataract formation. They may also be used to treat

CC neurodegenerative diseases, renal and kidney disease, bone and tooth

CC abnormalities, wounds and skin damage, e.g. by use of the nucleic acid in

CC gene therapy by using antibodies directed against CRIM1 polypeptides.

CC The present sequence represents C. elegans CRIM1 (AKA S52).

XX

SQ Sequence 872 AA;

Query Match 22.0%; Score 1307; DB 22; Length 872;

Best Local Similarity 32.9%; Pred. No. 1.3e-66;

Matches 280; Conservative 112; Mismatches 306; Indels 154; Gaps 36;

QY 173 DCSKARCEVQSPRCPEDSVLIEGYAPPBGCCPLPSRCVNCNAGCLRKV--CQPGMLNIL 230

Db 4 DCLKAICPLVFKHGKCPDSQLITVSPAPGNCCPPGSGCHDQKCKVPSVPTCTKEERLVM 63

QY 231 VSKASGKPGCCDLIECKPVGVDCRTVEQTV--OQTACPPDSYETQVRLTADGCTLP 288

Db 64 VEGSDIPGKCCAYECHKK-EKKCNVHCPPMFQDEECPPDSIRPPSISESCCPIR 122

QY 289 TRCELSGLGFPVCEVSGTPRIVSRGDGTFPGKCCDVFECVND--TKPACVFNNVYYDG 346

Db 123 QSCCKRGSIICRPAQC PDGKVNVTKGTGFPGRCCDKWECVDAELSKAKCNHSGIERQPL 182

QY 347 DMFRMNCRCRCGGVAICFTACGGEIN--CERYVPEGECPCVEDPVYPNPNAGCY 404

Db 183 ETHVSDCESQCQIRGVSVCKNMTCPKVNOECTWIGTIGTCECPVC-----LGCT 232

QY 405 ANGLILAHGDRWEDDCCTFCQCVN-GERHCVATVCGTCTNPVKVPGCECPVCEETIIT 463

Db 233 DNQTKLKGATWQKDDCTCTSELGAHMCCKYKCTDCENPRKVEGQCCPCVDEPIIR 292

QY 464 VDPPA-CGELSNCTLTRKDCINGFKRDHNGCRTCQCINTQELGSEKRGQCTLNCPPGFLT 522

Db 293 --PPATCPSLELSLR--CANGLRDNIICYVCECLPDEV----- 328

QY 523 DAQNCETCECRPRPKKCRPI---ICDKYCPGLGLLNKHGCDICRCKKCPPELSCKICPL 578

Db 329 -----PTNPRCRLNDENCEKQCAHGYLKDEGCTGVCCKSKCPPLHQCHKCLY 377

QY 579 GFQODSHGCLICKREAS-----ASAGPPILSGTCLTV--DGHH--HK 617

Db 378 GFETNSAGCSLCKCRASSKLDKQGTGTTKLGAGSAQTEYHSEKISFNSDGHQIVRD 437

QY 618 NEESWHDGCRECYCLNGREMCALITCPV-PA-CGNP--TIHPGCCPCSCADDFVQKPEL 673

Db 438 GGEWWSGCRHCFCEKQEFCSLSICTKPSDCADENKWKQKEDECCPSCIDQ--KKKPKS 495

QY 674 STP-----SICHAPG-GEYFVEGETWNIQDCTQCTCHSGRVLCETEVCPPLQNPS 724

Db 496 SNLSAAQKHEHTVCQSPGTGRFTDGTWQAPCVSCTCRVGHVLCRTTECPPIACPNPE 555

QY 725 -RTQDSCCPOCTDQPPRPSLRNNSVNYCKNDEGDIFL---AASWKPVDVCTSCIC-I 778

Db 556 YQNEEDCCPTCPEQ-----KVEN-TKNEKGDITVCTDDTAHIVDDTSCVCSA 604

QY 779 DSVTSCFSESC-PSVSCE-RPVLKKGCCPYCIKDTIPKKVCHGFSKAYADERWDLDS 836

Db 605 EGSADCYKEADESLECRGNPLVIKGCPCVC-SDALSSASVCSQSSVYAIQEQWDGR 663

QY 837 CTHCYCLOQ-GQTLGCTVSCPPPLPCVPEPINVEGSCCPMCPMEYVPEPTNPIEKTNRGEV 895

Db 664 CSNCSCTVGGTVCQRMVCP--HCCDDPVPIEGHCCPLCKD----- 701

QY 896 DLEVPLW-PTPSENDIVHLPRDMGHLOVDYDRNRLHPSDESSLDs---IASVVVPIIICLS 952
Db 702 -----AKSPYGYGNGSASFETSLG-----PRVDDGNGSSATSILIVVSLMSLCVVALI 749
QY 953 IIIAFLFINOKK 964
Db 750 IVLMLLYKRNRK 761

RESULT 14
AAM25238
ID AAM25238 standard; Protein; 193 AA.
XX AC AAM25238;
XX 16-OCT-2001 (first entry)
XX Human protein sequence SEQ ID NO:753.
XX
KW Human; cancer; ulcer; HIV infection; human immunodeficiency virus;
KW antiinflammatory; antirheumatic; antiarthritic; immunosuppressive;
KW antibacterial; endocrine; cardiant; central nervous system; virucide;
KW anti-HIV; fungicide; antimutagen; cardiovascular; antianaemic; anaemia;
KW antiaggregant; haemostatic; antidiabetic; antidiabetic; osteopathic; eczema;
KW dermatological; antiallergic; antiasthmatic; antiparkinsonian; infection;
KW neuroprotective; antidepressant; nootropic; antiparkinsonian; infection;
KW immunostimulant; gene therapy; antisense therapy; vaccine; inflammation;
KW antianaphylactic; rheumatoid arthritis; septic shock; pancreatitis;
KW cardiac dysfunction; neuropathology; cardiac anaphylaxis; autolmmunity;
KW genetic disease; haematopoietic disorder; platelet disorder; asthma;
KW thrombocytopaenia; osteoporosis; severe combined immunodeficiency;
KW allergic rhinitis; diabetes; multiple sclerosis; depression;
KW Alzheimer's disease; Parkinson's disease; neurodegenerative disorder;
KW neurological disorder.
XX Homo sapiens.
XX OS
XX PN WO200153455-A2.
XX PD 26-JUL-2001.
XX 22-DEC-2000; 2000WO-US35017.
XX PF
XX PR 23-DEC-1999; 99US-0471275.
XX PR 21-JAN-2000; 2000US-0488725.
XX PR 25-APR-2000; 2000US-0552317.
XX PA (HYSE-) HYSEQ INC.
XX PI Tang YT, Liu C, Drmanac RT;
XX WPI; 2001-457603/49.
XX N-PSDB; AAH99179.
XX
PT Isolated human polynucleotides encoding polypeptides, useful for the
PT treatment and diagnosis of e.g. cancer, ulcers and HIV infection -
XX
XX Claim 20; Page 181; 1217pp; English.
XX
XX AAH99166 to AAH99904 encode the human proteins given in AAM25225 to
XX AAM25963. The proteins can have activities based on the tissues and
XX cells they are expressed in, such as: antiinflammatory; antirheumatic;
XX antiarthritic; immunosuppressive; antibacterial; endocrine; cardiant;
XX central nervous system; virucide; anti-HIV; fungicide; antimutagen;
XX cardiovascular; antianaemic; antiaggregant; haemostatic; vulneryary;
XX antiulcer; osteopathic; dermatological; antiallergic; antiasthmatic;
XX antidiabetic; cytostatic; neuroprotective; antidepressant; nootropic;
XX antiparkinsonian; and immunostimulant. The proteins and polynucleotides
XX encoding them can be used in gene therapy, antisense therapy and vaccine
XX production. The proteins and polynucleotides are useful for screening for
XX agonists or antagonists of a protein and for the treatment and diagnosis
XX of disorders associated with the activity of a protein e.g. inflammation,
XX rheumatoid arthritis, septic shock, pancreatitis, cardiac dysfunction,

CC neuropathology, cardiac anaphylaxis, viral, bacterial, HIV and fungal
CC infections, autoimmunity, genetic diseases, haematopoietic disorders,
CC anaemia, platelet disorders, thrombocytopaenia, wounds, burns, ulcers,
CC osteoporosis, severe combined immunodeficiency, eczema, allergic
CC rhinitis, asthma, diabetes, cancer, multiple sclerosis, depression,
CC Alzheimer's disease, Parkinson's disease, neurodegenerative and
CC neurological disorders.
XX
SQ Sequence 193 AA;
Query Match 17.5%; Score 1042; DB 22; Length 193;
Best Local Similarity 98.4%; Pred. No. 3.7e-52;
Matches 188; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 10 LAGCGHLLVSLGLLLPARSGTRALVCLPCDESKCEPRNRPGSIYQVCGCGCYTCASQ 69
Db 3 LAGCGHLLVSLGLLLPARSGTRALVCLPCDESKCEPRNRPGSIYQVCGCGCYTCASQ 62
QY 70 GNECGGTFGIYGTCDRLGRCVIRPPLNGDSLTYEAGVCEDEENWTDQLLGFKPCNENL 129
Db 63 RNESCGGTFGIYGTCDRLGRCVIRPPLNGDSLTYEAGVCEDEENWTDQLLGFKPCNENL 122
QY 130 IAGCNIINGKCECWTIRTCNPFEPFQSDMCLSKALKRIEKEKPCDKARCEVQFSPRCE 189
Db 123 IAGCNIINGKCECWTIRTCNPFEPFQSDMCLSKALKRIEKEKPCDKARCEVQFSPRCE 182
QY 190 DSVLIIEGYAPP 200
Db 183 DSVLIIEGYAPP 193
RESULT 15
AAU83112
ID AAU83112 standard; Protein; 225 AA.
XX AC AAU83112;
XX DT 08-MAY-2002 (first entry)
XX DE Novel secreted protein Z790708GIP.
XX KW Protein secretion; mammalian secreted polypeptide; MSP.
XX OS Homo sapiens.
XX PN WO200202621-A2.
XX PD 10-JAN-2002.
XX PF 28-JUN-2001; 2001WO-US20638.
XX PR 30-JUN-2000; 2000US-215446P.
XX PA (ZYMO) ZYMOGENETICS INC.
XX PI Sheppard PO, Presnell SR;
XX WPI; 2002-147999/19.
XX N-PSDB; ABK33027.
XX
PT Novel isolated mammalian secreted polypeptide useful in therapeutic and
PT diagnostic methods, to direct secretion of other proteins of interest
PT from host cell, as educational tools, and as laboratory practicum kits
XX
XX Claim 12; Page 135; 397pp; English.
XX The invention describes an isolated mammalian secreted polypeptide (MSP)
XX (I). (I) is useful to direct the secretion of other proteins of interest
XX from a host cell, to monitor secretion of proteins, to degenerate
XX sequences comprising all nucleotide sequences encoding a particular
XX polypeptide, to screen for cell metabolism effecting receptors, for
XX identifying new target receptors and drug design, for identifying, for

CC protein purification, for determining the weight of expressed MSP
CC polypeptides as a ratio to total protein expressed, for identifying
CC peptide cleavage sites, for coupling amino and carboxy terminal tags, for
CC amino acid sequence analysis, for monitoring biological activities of the
CC protein in vitro and in vivo, and to teach analytical skills and as
CC reagents for the study of cells, receptors, and other binding molecules.
CC The polynucleotide is useful for radiation hybrid mapping, and somatic
CC cell genetic technique developed for constructing high-resolution,
CC contiguous maps of mammalian chromosomes. Reagents disclosed in the
CC invention may be used to detect metabolic abnormalities characterised by
CC over or under production of the protein. This is the amino acid sequence
CC of a mammalian secreted polypeptide, described in the method of the
CC invention.
xx

SQ Sequence 225 AA;

Query Match 9.8%; Score 582; DB 23; Length 225;
Best Local Similarity 97.3%; Pred. No. 7.1e-26;
Matches 107; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 MYLVAGDRGLACGCHLLVSLGLLLPARSGTRALVCLPCDESKCEEPNRPGSIVOGVC 60

Db 1 MYLVAGDRGLACGCHLLVSLGLLLPARSGTRALVCLPCDESKCEEPNRPGSIVOGVC 60

QY 61 GCCYTCA SQNESCGGTGFIYGTCDRGLRCVIRPPLNGDSLTYEAGVCE 110

Db 61 GCCYTCA SQNESCGGTGFIYGTCDRGLRCVIRPPLNGDSLTYEAGVCE 110

Search completed: March 14, 2003, 17:40:36
Job time : 50 secs

